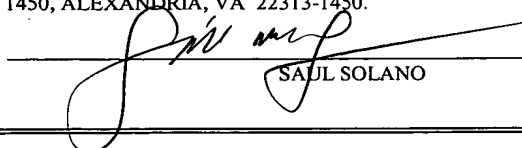


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SAUL SOLANO

**CONTINUATION-IN-PART
APPLICATION**

for

UNITED STATES LETTERS PATENT

on

**TETRACALCIUM PHOSPHATE (TTCP) HAVING CALCIUM PHOSPHATE
WHISKER ON SURFACE AND PROCESS FOR PREPARING THE SAME**

by

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**TETRACALCIUM PHOSPHATE (TTCP) HAVING CALCIUM PHOSPHATE
WHISKER ON SURFACE AND PROCESS FOR PREPARING THE SAME**

CROSS-REFERENCE TO RELATED APPLICATIONS

[0001] The present application is a continuation-in-part application of US Patent Application Serial Number 10/607,023, filed June 27, 2003, which is a continuation-in-part application of U.S. Patent Application Serial Number 10/414,582, filed April 16, 2003, which is a continuation-in-part application of U.S. Patent Application Serial Number 09/615,384, filed July 13, 2000, now abandoned, which is a continuation-in-part application of U.S. Patent Application Serial Number 09/351,912, filed July 14, 1999, now US patent No. 6,379,453B1. The above-listed applications are commonly assigned with the present invention and the entire contents of which are incorporated herein by reference.

BACKGROUND OF THE INVENTION

FIELD OF THE INVENTION

[0002] The present invention relates to a tetracalcium phosphate (TTCP) for producing fast-setting, bioresorbable calcium phosphate cements (CPC), and in particular, to a tetracalcium phosphate having whiskers on the surface thereof for producing fast-setting, bioresorbable CPC having a high initial strength.

DESCRIPTION OF THE RELATED ART

[0003] U.S. Pat. No. 6,379,453B1 which is commonly assigned with the present invention discloses a process for producing a fast-setting, bioresorbable calcium phosphate cement comprising the following steps: obtaining a powder mixture from at least one calcium phosphate selected from the group consisting of $\text{Ca}_4(\text{PO}_4)_2\text{O}$, $\text{CaHPO}_4 \cdot 2\text{H}_2\text{O}$, CaHPO_4 , $\text{Ca}_8\text{H}_2(\text{PO}_4)_6 \cdot 5\text{H}_2\text{O}$, $\alpha\text{-Ca}_3(\text{PO}_4)_2$, $\beta\text{-Ca}_3(\text{PO}_4)_2$, $\text{Ca}_2\text{P}_2\text{O}_7$, $\text{Ca}_2\text{H}_2\text{P}_2\text{O}_8$, wherein the molar ratio of Ca to P in the mixture is roughly between 1 and 2; mixing the powder mixture in a phosphate-containing solution to obtain a powder/solution mixture

having a concentration of less than 4 g powder mixture per ml solution; immediately heating the powder/solution mixture to a temperature of roughly 50°C-350°C to obtain a powder containing uniformly distributed submicron-sized apatite crystals; and mixing the apatite crystal-containing powder in a phosphate ion-containing solution to obtain a fast-setting, bioresorbable calcium phosphate cement.

SUMMARY OF THE INVENTION

[0004] An extensive study on the preparation of the fast-setting, bioresorbable calcium phosphate cement disclosed in U.S. Pat. No. 6,379,453B1 has been conducted by the same inventors and their co-workers, and found that a fast-setting, bioresorbable CPC having a high initial strength can be prepared from a unique calcium phosphate, tetracalcium phosphate ($\text{Ca}_4(\text{PO}_4)_2\text{O}$, TTCP) particle having basic whiskers or fine crystals on the surface thereof, wherein said basic whiskers or fine crystals have a Ca/P ratio greater than 1.33. Therefore an object of the invention is to provide such a unique TTCP particle. Another object of the present invention is to provide a process for preparing said unique TTCP particle. A further object of the present invention is to provide a fast-setting, bioresorbable CPC calcium phosphate cement prepared from said unique TTCP particle.

[0005] The invention accomplishes the above object by providing a tetracalcium phosphate ($\text{Ca}_4(\text{PO}_4)_2\text{O}$, TTCP) particle having basic calcium phosphate whiskers on a surface of said TTCP particle; said basic calcium phosphate whiskers having a length up to about 5000 nm and a width up to about 500 nm, and preferably, a length from about 1 nm to about 2000 nm and a width from about 1 nm to about 200 nm. Said basic calcium phosphate whiskers have a Ca/P molar ratio greater than 1.33, and preferably greater than 1.35 and less than 4.0. Said basic calcium phosphate whiskers have a non-stoichiometric chemical composition. Further, said basic calcium phosphate whiskers are substantially free of a hydroxyapatite phase, and comprises TTCP as a major phase.

BRIEF DESCRIPTION OF THE DRAWINGS

[0006] Figs. 1A to 1C are related to microstructure and diffraction pattern of calcium phosphate whiskers grown on TTCP surface according to the present invention, wherein (a) bright field image of whiskers; (b) electron diffraction pattern of whiskers; and (c) interpretation of the diffraction pattern.

[0007] Fig. 2 shows XRD patterns, wherein (a) TTCP without whisker treatment; (b) TTCP with whisker treatment in $(\text{NH}_4)_2\text{HPO}_4$ for 5 minutes; and (c) CPC prepared from whisker-treated TTCP powder immersed in Hanks' solution for 24 hours.

DETAILED DESCRIPTION OF THE INVENTION

[0008] The present invention discloses a process for preparing a tetracalcium phosphate (TTCP) powder comprising TTCP particles comprising basic calcium phosphate whiskers on surfaces of said TTCP particles, said process comprising the following steps:

a) mixing a TTCP powder with a whisker-inducing solution so that basic calcium phosphate whiskers start to grow on surfaces of TTCP particles of said TTCP powder;

b) terminating the growth of said calcium phosphate whiskers by drying the whisker-inducing solution in the mixture, so that said calcium phosphate whiskers have a length up to about 5000 nm and a width up to about 500 nm, and preferably, a length from about 1 nm to about 2000 nm and a width from about 1 nm to about 200 nm, said basic calcium phosphate whiskers have a Ca/P molar ratio greater than 1.33, preferably greater than 1.35 and less than 4.0, and said basic calcium phosphate whiskers have a non-stoichiometric chemical composition, preferably said basic calcium phosphate whiskers are substantially free of a hydroxyapatite phase, and comprises TTCP as a major phase.

[0009] Optionally, at least one additive selected from the group consisting of sodium phosphate (Na_3PO_4), disodium hydrogen phosphate (Na_2HPO_4), sodium dihydrogen phosphate (NaH_2PO_4), disodium hydrogen phosphate dodecahydrate ($\text{Na}_2\text{HPO}_4 \cdot 12\text{H}_2\text{O}$),

disodium hydrogen phosphate heptahydrate ($\text{Na}_2\text{HPO}_4 \cdot 7\text{H}_2\text{O}$), sodium phosphate dodecahydrate ($\text{Na}_3\text{PO}_4 \cdot 12\text{H}_2\text{O}$), orthophosphoric acid (H_3PO_4), calcium sulfate (CaSO_4), $\text{Ca}_4(\text{PO}_4)_2\text{O}$, $\text{CaHPO}_4 \cdot 2\text{H}_2\text{O}$, CaHPO_4 , $\text{Ca}_8\text{H}_2(\text{PO}_4)_6 \cdot 5\text{H}_2\text{O}$, $\alpha\text{-Ca}_3(\text{PO}_4)_2$, $\beta\text{-Ca}_3(\text{PO}_4)_2$, $\text{Ca}_2\text{P}_2\text{O}_7$, and $\text{Ca}_2\text{H}_2\text{P}_2\text{O}_8$, $(\text{NH}_4)_3\text{PO}_4$, $(\text{NH}_4)_2\text{HPO}_4$, and $(\text{NH}_4)\text{H}_2\text{PO}_4$ together with said TTCP particles are mixed with the whisker-inducing solution in step a).

[0010] Optionally, said drying in step b) is carried out by heating the mixture resulting from step a) at a temperature less than about 1000°C . Preferably, said drying in step b) comprises separating the mixture resulting from step a), and heating the separated powder at a temperature of about 50 to 500°C .

[0011] The heating includes (but not limited to) the conventional oven/furnace heating, resistance heating, infrared heating, microwave heating, electron beam heating, ion beam heating, laser beam heating and plasma heating. Preferably said heating is conducted in vacuum, inert atmosphere or air atmosphere.

[0012] The whisker-inducing solution in step a) may be an acidic aqueous solution, a basic aqueous solution, an organic solvent or a substantially pure water. The acidic aqueous solution may contain at least one Ca or P source, or is free from Ca and P. The acidic aqueous solution can be selected from the group consisting of nitric acid (HNO_3), hydrochloric acid (HCl), phosphoric acid (H_3PO_4), carbonic acid (H_2CO_3), sodium dihydrogen phosphate (NaH_2PO_4), sodium dihydrogen phosphate monohydrate, sodium dihydrogen phosphate dihydrate, potassium dihydrogen phosphate (KH_2PO_4), ammonium dihydrogen phosphate ($\text{NH}_4\text{H}_2\text{PO}_4$), malic acid, acetic acid, lactic acid, citric acid, malonic acid, succinic acid, glutaric acid, tartaric acid, oxalic acid and their mixture.

[0013] The basic aqueous solution for use as the whisker-inducing solution in the method of the present invention may contain at least one Ca or P source, or is substantially free from Ca and P. The basic aqueous solution may be selected from the group consisting of ammonia, ammonium hydroxide, alkali metal hydroxide, alkali earth hydroxide, disodium hydrogen phosphate (Na_2HPO_4), disodium hydrogen phosphate dodecahydrate, disodium hydrogen phosphate heptahydrate, sodium phosphate dodecahydrate ($\text{Na}_3\text{PO}_4 \cdot 12\text{H}_2\text{O}$), dipotassium hydrogen phosphate (K_2HPO_4), potassium

phosphate tribasic (K_3PO_4), diammonium hydrogen phosphate ($(NH_4)_2HPO_4$), ammonium phosphate trihydrate ($(NH_4)_3PO_4 \cdot 3H_2O$), sodium bicarbonate ($NaHCO_3$), and their mixture.

[0014] Preferably, said whisker-inducing solution in step a) is a basic aqueous solution. More preferably, said basic aqueous solution is a diammonium hydrogen phosphate ($(NH_4)_2HPO_4$), Na_2HPO_4 , or K_2HPO_4 aqueous solution. A suitable diammonium hydrogen phosphate ($(NH_4)_2HPO_4$) aqueous solution has a concentration of at least 5 wt%, preferably 10-60 wt%, based on the weight of said solution, and the mixing of said TTCP powder with this diammonium hydrogen phosphate ($(NH_4)_2HPO_4$) aqueous solution in step a) is in a ratio of less than about 10 g powder per ml solution, preferably less than about 5 g powder per ml solution. In one of the preferred embodiment of the present invention, said concentration is about 33 wt%, and the mixing ratio is about 1 gm TTCP per 13 ml solution.

[0015] The present invention also discloses a calcium phosphate cement (CPC) powder comprising the TTCP powder of the present invention.

[0016] The following examples are intended to demonstrate the invention more fully without acting as a limitation upon its scope, since numerous modifications and variations will be apparent to those skilled in this art.

TTCP preparation

[0017] The TTCP powder was fabricated in-house from the reaction of dicalcium pyrophosphate ($Ca_2P_2O_7$) (Sigma Chem. Co., St. Louis, MO, USA) and calcium carbonate ($CaCO_3$) (Katayama Chem. Co., Tokyo, Japan) using the method suggested by Brown and Epstein [*Journal of Research of the National Bureau of Standards- A Physics and Chemistry* 6 (1965) 69A 12].

TEM examination

[0018] A Hitachi Model-HF2000 200kV field emission transmission electron microscope (TEM) equipped with a Noran Vayager Model 1000 energy dispersive

spectroscopy (EDS) system was used for the study. The aperture size for microchemical analysis (Ca/P ratio) is 15nm.

EXAMPLE 1

WHISKER-INDUCING TREATMENT OF TTCP PARTICLES TREATED IN PHOSPHATE-CONTAINING BASIC SOLUTION

[0019] $\text{Ca}_4(\text{PO}_4)_2\text{O}$ (TTCP) powder as synthesized was sieved with a #325 mesh. The sieved powder has an average particle size of about 10 μm . An aqueous solution of diammonium hydrogen phosphate was prepared by dissolving 20 g of diammonium hydrogen phosphate, $(\text{NH}_4)_2\text{HPO}_4$, in 40 ml deionized water. The resulting solution had a pH value of 8.02. To the TTCP powder the basic aqueous solution of diammonium hydrogen phosphate was added according to the ratio of 1 gm TTCP/13 ml solution. The TTCP powder was immersed in the basic aqueous solution for various periods of time of 1 minute, 5 minutes and 10 minutes, and filtered rapidly with a vacuum pump again. The resulting powder cake was dried in an oven at 50°C. The dried powder was dispersed in ethanol with supersonication. A drop of the dispersion was dripped on a single-side carbon sieve of #325 mesh having a diameter of 3 mm, and left dry to obtain a specimen coated with a thin carbon film for electrical conductivity for TEM examination. The microchemical analysis (Ca/P ratio) results of ten specimens (P1 to P10) for each treat time are shown in Table 1.

Table 1

Treat time	Ca/P											SD*	Whisker width (nm)	Whisker length (nm)
	P1	P2	P3	P4	P5	P6	P7	P8	P9	P10	Avg.			
1 min	1.20	1.30	1.26	1.14	1.12	1.03	1.22	1.19	1.14	1.25	1.19	0.08	<50	<100
5 min	1.85	1.61	1.35	1.76	1.40	1.52	1.63	1.53	1.35	1.38	1.54	0.17	<100	<300
10 min	3.81	3.20	1.78	1.74	1.80	1.38	1.61	1.81	2.01	1.63	2.08	0.78	<100	<300

*SD= standard deviation

[0020] FIG. 1 represents a typical microstructure of the calcium phosphate whiskers grown on TTCP surface under such condition. FIG. 1A is a bright-field image showing the whiskers are substantially radial-oriented and the majority of which have lengths <300nm and widths <100nm; FIG. 1B is a typical electron diffraction pattern of such whiskers. The dotted-ring pattern is a direct result of the diffraction of numerous nano-sized whiskers; FIG. 1C is the indexing/interpretation of the diffraction pattern, which clearly shows that every ring matches a certain crystallographic plane of TTCP phase, indicating the whiskers have a TTCP crystal structure. The absence of hydroxyapatite (HA) phase (100) ring ($d = 0.817 \text{ nm}$) in the diffraction pattern excludes the possibility for the whiskers to have an apatite crystal structure under this whisker treatment condition. It also can be seen from Table 1 that basic calcium phosphate whiskers have a Ca/P ratio other than 1.67, i.e. a non-stoichiometric chemical composition. The Ca/P ratio of hydroxyapatite (HA) is 1.67. The results show that Ca/P ratio is sensitive to the process condition (in this case, treating time).

EXAMPLE 2

WHISKER-INDUCING TREATMENT OF TTCP PARTICLES TREATED IN PHOSPHATE-CONTAINING ACIDIC SOLUTION

[0021] The procedures of Example 1 were repeated except that the basic aqueous solution was changed to 1M phosphorus acid aqueous solution having a pH of 0.8 and the immersion time was changed to 30 seconds. The results are shown in Table 2.

Table 2

Treat time	Ca/P											SD*	Whisker width (nm)	Whisker length (nm)
	P1	P2	P3	P4	P5	P6	P7	P8	P9	P10	Avg.			
30 sec	3.73	2.0	2.28	1.41	2.65	1.43	1.77	1.89	1.65	1.54	2.04	0.71	<200	<600

*SD= standard deviation

EXAMPLE 3
WHISKER-INDUCING TREATMENT OF TTCP PARTICLES TREATED IN
PHOSPHATE-FREE BASIC SOLUTION

[0022] The procedures of Example 1 were repeated except that the basic aqueous solution was changed to a basic aqueous NaOH solution having a pH of 10.66 and the immersion time was changed to 30 seconds and 24 hours. For the specimens treated for 30 seconds no whisker was observed on TTCP surface. The results for the treat time of 24 hours are shown in Table 3.

Table 3

Treat time	Ca/P											SD*	Whisker width (nm)	Whisker length (nm)
	P1	P2	P3	P4	P5	P6	P7	P8	P9	P10	Avg.			
24 hr	1.90	2.19	2.80	3.40	1.47	2.05	1.53	1.63	1.42	2.03	2.04	0.63	<200	<600

*SD= standard deviation

EXAMPLE 4
WHISKER-INDUCING TREATMENT OF TTCP PARTICLES TREATED IN
PHOSPHATE-FREE ACIDIC SOLUTION

[0023] The procedures of Example 1 were repeated except that the basic aqueous solution was changed to 0.16M HCl aqueous solution having a pH of 0.8 and the immersion time was changed to 30 seconds, 10 minutes, one hour and 24 hours. For the specimens treated for 30 seconds no whisker was observed on TTCP surface. The results for the remaining treat times are shown in Table 4.

Table 4

Treat time	Ca/P											SD*	Whisker width (nm)	Whisker length (nm)
	P1	P2	P3	P4	P5	P6	P7	P8	P9	P10	Avg.			
10 min	1.61	1.92	1.40	1.52	1.52						1.59	0.20	<50	<100
1 hr	1.41	1.90	1.52	1.67	1.57	1.42	1.53	1.46	1.38	1.60	1.55	0.15	<100	<200
24 hr	2.65	1.53	1.61	1.77	1.52	2.23	1.36	1.83	1.44	2.09	1.80	0.41	<200	<600

*SD= standard deviation

EXAMPLE 5

COMPRESSIVE STRENGTH OF CPC PREPARED FROM THE WHISKER-GROWN TTCP PARTICLES

[0024] $\text{Ca}_4(\text{PO}_4)_2\text{O}$ (TTCP) powder as synthesized was sieved with a #325 mesh and has an average particle size of about 10 μm . To the sieved TTCP powder a HCl aqueous solution having a pH of 0.8 was added according to the ratio of 1 gm TTCP/13ml solution. The sieved TTCP powder was immersed in the HCl solution for 12 hours, filtered rapidly and washed with deionized water, and filtered rapidly with a vacuum pump again. The resulting powder cake was dried in an oven at 50°C. The dried powder was divided into halves, ground for 20 minutes and 120 minutes separately, and combined. A setting solution of diammonium hydrogen phosphate was prepared by dissolving 20 g of diammonium hydrogen phosphate, $(\text{NH}_4)_2\text{HPO}_4$, in 40 ml deionized water. 100 g of the mixed ground powder and 35 ml of the setting solution were well mixed to form a paste, which was then filled in molds to form specimens for compression test. The specimens were removed from the molds 15 minutes after the mixing, and soaked in a Hanks' solution. The soaked specimens were removed from the Hanks' solution at various periods of soaking time, and were immediately subjected to the compression test without drying. The compression test was conducted according to a method commonly used in the literature. The cylindrical samples have a diameter of 6 mm and a length of 12 mm. Results: compressive strength is 27.4 MPa for the soaking time of 20 minutes, and 48 MPa for one-day soaking time.

EXAMPLE 6
COMPRESSIVE STRENGTH OF CPC PREPARED FROM THE
WHISKER-GROWN TTCP PARTICLES

[0025] $\text{Ca}_4(\text{PO}_4)_2\text{O}$ (TTCP) powder as synthesized was sieved with a #325 mesh and has an average particle size of about 10 μm . To the sieved TTCP powder the aqueous $(\text{NH}_4)_2\text{HPO}_4$ solution prepared in Example 1 was added according to the ratio of 1 gm TTCP/13ml solution. The sieved TTCP powder was immersed in the $(\text{NH}_4)_2\text{HPO}_4$ solution for 5 minutes, filtered rapidly and washed with deionized water, and filtered rapidly with a vacuum pump again. The resulting powder cake was dried in an oven at 50°C. The dried powder was ground 120 minutes to obtain a powder A. The procedures in Example 5 were repeated to obtain a powder B except that the dried powder was ground only for a period of 300 minutes. A mixed powder of A and B in a ratio of 1:1 ratio was subjected to the compression tests following the procedures recited in Example 5. Results: compressive strength is 26 MPa for the soaking time of 20 minutes, and 42.8 MPa for one-day soaking time.

EXAMPLE 7
COMPRESSIVE STRENGTH OF CPC PREPARED FROM THE
WHISKER-GROWN TTCP PARTICLES

[0026] The procedures in Example 5 were repeated except that the HCl solution was changed to the aqueous $(\text{NH}_4)_2\text{HPO}_4$ solution prepared in Example 1 and the soaking time was changed to 5 minutes. Results: compressive strength is 18.6 MPa for the soaking time of 20 minutes, and 48.8 MPa for one-day soaking time.

EXAMPLE 8
COMPRESSIVE STRENGTH OF CPC PREPARED FROM THE
WHISKER-GROWN TTCP PARTICLES

[0027] $\text{Ca}_4(\text{PO}_4)_2\text{O}$ (TTCP) powder as synthesized was sieved with a #325 mesh and ground for two hours. To the ground TTCP powder the powder B prepared in Example 5 was added and mixed in a ratio of 1:1. The resulting mixed powder was subjected to the compression tests following the procedures recited in Example 5. Results: compressive strength is 19.7 MPa for the soaking time of 20 minutes, and 43.6 MPa for one-day soaking time.

EXAMPLE 9
X-RAY DIFFRACTION OF WHISKER-TREATED TTCP POWDER AND
IMMERSED CPC PREPARED FROM SUCH TTCP

[0028] A TTCP powder was whisker-treated for 5 minutes according to the process described in Example 1. X-ray diffraction (XRD) was performed using an X-ray diffractometer (Rigaku D-max IIIV, Tokyo, Japan) with Ni-filtered $\text{CuK}\alpha$ radiation operated at 30 kV and 20 mA at a scanning speed of $1^\circ/\text{min}$. The phases were identified by matching each characteristic XRD peak with that compiled in JCPDS files.

[0029] Results: As indicated in Fig. 2, the XRD pattern of the whisker-treated TTCP powder (b) is substantially identical to that of TTCP as synthesized (a). The perfect match of every XRD peak position (diffraction angle) with the JCPDS data indicates that there is no additional phase formed during the whisker treatment. 0.7 g whisker-treated TTCP powder with 0.25 ml setting solution to form a CPC paste. The setting solution was prepared by dissolving 20 g $(\text{NH}_4)_2\text{HPO}_4$ in 40 ml deionized water. The CPC paste was filled in a cylindrical mold (12 mm in height and 6 mm in diameter), allowing hardening of the paste to occur within the mold. After 15 minutes the hardened CPC sample was removed from the mold and immersed in a 37°C Hanks' solution for 24 hours. After removing from the Hanks' solution and drying, the CPC sample was ready for XRD

analysis. After immersion in Hanks' solution for 24 hours, the XRD pattern (c) of the CPC shows a large amount of HA phase which has replaced TTCP as the dominant phase. At this time only a small amount of TTCP remains. The result suggests that the CPC prepared from the whisker-treated TTCP powder of the invention can quickly transform into HA (the major component of human bone), once implanted.

EXAMPLE 10
SETTING SOLUTION PREPARED FROM $(\text{NH}_4)\text{H}_2\text{PO}_4$ AND KOH

[0030] A TTCP powder was whisker-treated for 5 minutes according to the process described in Example 1. The resulting powder cake was dried in an oven at 50°C. The dried powder was ground for 120 minutes. A setting solution was prepared by dissolving 13.2 g $(\text{NH}_4)\text{H}_2\text{PO}_4$ in 40 ml deionized water to obtain an initial solution having a pH value of 3.72, and adding KOH to the initial solution so that the pH value was adjusted to 7.5. 100 g of the ground powder and 35 ml of the setting solution were well mixed to form a paste for 1 minute, which was then filled in molds to form specimens for compression tests following the procedures recited in Example 5. Results: compressive strength is 9.6 MPa for the soaking time of 20 min.

EXAMPLE 11
SETTING SOLUTION PREPARED FROM $(\text{NH}_4)\text{H}_2\text{PO}_4$ AND NaOH

[0031] The procedures in Example 10 were repeated except that the KOH was changed to NaOH and the final pH value of the setting solution was 7.8, and 20 ml of the setting solution was mixed with 100 g of the ground powder. Results: compressive strength is 10.3 MPa for the soaking time of 20 min.

EXAMPLE 12

**SETTING SOLUTION PREPARED FROM $(\text{NH}_4)_2\text{HPO}_4$, $\text{NaH}_2\text{PO}_4 \cdot 2\text{H}_2\text{O}$
AND K_2HPO_4**

[0032] A TTCP powder was prepared following the procedures recited in Example 10. A setting solution was prepared by dissolving 7.5 g $(\text{NH}_4)_2\text{HPO}_4$, 2.5 g $\text{NaH}_2\text{PO}_4 \cdot 2\text{H}_2\text{O}$ and 5 g K_2HPO_4 in 40ml deionized water. The final pH value of the setting solution was 7.56. 100 g of the ground powder and 30 ml of the setting solution were well mixed to form a paste for 1 minute, which was then filled in molds to form specimens for compression tests following the procedures recited in Example 5. Results: compressive strength is 18.0 MPa for the soaking time of 20 min.

EXAMPLE 13

**SETTING SOLUTION PREPARED FROM $\text{Na}_2\text{HPO}_4 \cdot 12\text{H}_2\text{O}$, $\text{NaH}_2\text{PO}_4 \cdot 2\text{H}_2\text{O}$
AND $(\text{NH}_4)_2\text{HPO}_4$**

[0033] A TTCP powder was prepared following the procedures recited in Example 10. A setting solution was prepared by dissolving 3 g $\text{Na}_2\text{HPO}_4 \cdot 12\text{H}_2\text{O}$, 3 g $\text{NaH}_2\text{PO}_4 \cdot 2\text{H}_2\text{O}$ and 7.5g $(\text{NH}_4)_2\text{HPO}_4$ in 40ml deionized water. The final pH value of the setting solution was 7.38. 100 g of the ground powder and 30 ml of the setting solution were well mixed to form a paste for 1 minute, which was then filled in molds to form specimens for compression tests following the procedures recited in Example 5. Results: compressive strength is 20.8 MPa for the soaking time of 20 min.

EXAMPLE 14
SETTING SOLUTION PREPARED FROM PHOSPHORIC ACID AND
AMMONIA SOLUTION

[0034] A TTCP powder was prepared following the procedures recited in Example 10. A setting solution was prepared by mixing 37.68 ml of 85 wt% phosphoric acid and 100 ml deionized water, and then 73.8 ml of 28 wt% ammonia solution. The final pH value of the setting solution was 7.0. 100 g of the ground powder and 30 ml of the setting solution were well mixed to form a paste for 1 minute, which was then filled in molds to form specimens for compression tests following the procedures recited in Example 5. Results: compressive strength is 23.4 MPa for the soaking time of 20 min.

[0035] Although a “basic” whisker can be grown on TTCP surface by immersion in a variety of solutions, the process should be carefully controlled. For example, when the solution contains a P source in the absence of Ca, the immersion time should be long enough to grow a basic whisker (an “acidic” whisker is grown at the early stage due to the excess P ions in the solution). Yet the immersion time should not be too long either to avoid the basic whisker’s growing too large, that can largely deteriorate the CPC properties.

[0036] On the other hand, when the solution does not contain P (e.g., HCl), acidic whisker is never grown on the surface of TTCP particles. All the observed whiskers on TTCP particles at all stages are basic in nature.

[0037] In addition to Ca/P ratio, the growth rate of a basic whisker is also sensitive to such process parameters as the type, pH, temperature and ion concentrations of the solution, to name a few.

[0038] Although the present invention has been described with reference to specific details of certain embodiments thereof, it is not intended that such details should be regarded as limitations upon the scope of the invention except as and to the extent that they are included in the accompanying claims. Many modifications and variations are possible in light of the above disclosure.